PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference P-INCI-X-04-0273 International application No. PCT/EP2004/008510			FOR FURTHER A	√416					
			International filing date 29.07.2004	(day/month/year)	Priority date (day/month/year) 30.07.2003				
		• •	national classification and 409/12, CO7D513/04,		P3/04				
	licant BORATORIOS D	EL DR. ESTE	/E S.A. et al.						
1.			eliminary examination r			minary Examining			
2.	This REPORT co	onsists of a total	of 6 sheets, including	this cover sheet.					
3.	This report is als	o accompanied	by ANNEXES, compris	ing:					
	a. 🗆 sent to th	the applicant and to the International Bureau) a total of sheets, as follows:							
	and/c	ts of the descript or sheets contain nistrative Instruc	ion, claims and/or draw ing rectifications author tions).	ings which have bee ized by this Authorit	en amended and are the y (see Rule 70.16 and s	e basis of this report Section 607 of the			
sheets which supersede earlier sheets, but which this Authority considers contain an amendment that of beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.									
	sequence	listing and/or tal	Bureau only) a total of (bles related thereto, in Listing (see Section 8)	computer readable for	orm only, as indicated i	er(s)) , containing a n the Supplemental			
4.	This report conta	ins indications re	elating to the following	tems:					
☑ Box No. I Basis of the opinion									
	Box No. II Priority								
	☐ Box No. III	Non-establishm	ent of opinion with reg	ard to novelty, invent	tive step and industrial	applicability			
	☐ Box No. IV	Lack of unity of invention							
	Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement								
	☐ Box No. VI	Certain docume	ents cited						
			in the international app						
	☐ Box No. VIII	Certain observa	ations on the internation	nal application					
Date	of submission of the	demand		Date of completion of	of this report				
28.02.2005			30.06.2005						
Name and mailing address of the international				Authorized Officer					
preliminary examining authority: European Patent Office D-80298 Munich			Telephone No. +49	89 2399-	Statement Language . Elifa				
	Fax: +49 89	2399 - 0 Tx: 5236 9 2399 - 4465	556 epmu d			Tours outper on to be a series			

10/566 101 IAP9 Rec'd PCT/PTO 27 JAN 2006¹ International application No.

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	_	Box No. I	Basis of tl	ne report								
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)		Description	n, Pages									
		1-66			as originally fi	led						
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	3.		nendments h		ed in the ca	ncellatio	on of:					
			description, claims, Nos.									
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)	4.	had not bee	eport has bee en made, sin Ital Box (Rule	ce they ha								
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		* If it	em 4 appl:	ies, som	e or all	of th	ese she	eets ma	ay be ma	rked "	supersed	led."

Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

1-73

No:

Claims

1, 6-9

Inventive step (IS)

Yes: Claims

2-24

No:

Claims

1-73

Industrial applicability (IA)

Yes: Claims

1-73

No: Claims

2. Citations and explanations (Rule 70.7):

see separate sheet

Re Item V

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Reference is made to the following documents:

- D1: WO 03/042175 A (ESTEVE LABOR DR) 22 May 2003 (2003-05-22)
- D2: EP-A-0 815 961 (WABAG WASSERTECHN ANLAGEN GMBH) 7 January 1998 (1998-01-07)
- D3: WO 02/060871 A (FLAUGH MICHAEL EDWARD; GILLIG JAMES RONALD (US); HEINZ LAWRENCE JOSEP) 8 August 2002 (2002-08-08)
- D4: US-A-3 472 870 (GOULD BARBARA E ET AL) 14 October 1969 (1969-10-14)
- D5: BROWN F J ET AL: "Evolution of a Series of Peptidoleukotriene Antagonists: Synthesis and Structure-Activity Relationships of 1,6-Disubstituted Indoles and Indazoles" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY. WASHINGTON, US, vol. 6, no. 33, 1990, pages 1771-1781, XP002077393 ISSN: 0022-2623

2. Novelty (Article 33(1) and (2)PCT)

The present application discloses compounds of formula (la-b) (see present Claims 1, 9), which are useful as 5-HT-6 modulators.

The present compounds (see Claims 1 and 9) differ form the D1-D2 compounds on the account of the position 6 of substitution with sulfonamide moiety on the indole ring (instead of the 5-th position for the D1 compounds and 4-th position for the D2 compounds), from the D3 compounds on the account of the sulfonamide moiety (see Claim 1) and from the D4 compounds on the account of the A substituent of the sulfonamide function (which in the present case should contain a (hetero)aromatic ring and in D4 case is an alkyl or alkenyl chain). The D5 compounds differ from the present compounds on the account the N-benzyl moiety from the 1-th position of the indole ring (the present R1 substituent cannot be an aryl moiety). Consequently, the novelty of the present subject-matter is acknowledged.

3. Inventive step (Article 33(1) and 33(3) PCT)

The present application discloses 6-sulfonamidoindoles which are substituted on position 1 of the indole with an amino moiety or with a (hetero) cycloaliphatic ring useful as 5-HT-6 modulators.

D1, which is regarded as being the closest prior art, discloses 5-HT-6 modulators which are also sulfonamide indoles. The main differences between the present compounds and the compounds disclosed by D1 are the positions of substitution with the sulfonamide moiety and -(CH2)n-R1 substituent on the indole ring (see present Claim 1 and the Claim 1 of D1).

The problem underlying the present invention cannot be regarded in providing further sulfonamide indoles useful as to treat diseases modulated through the 5-HT-6 receptor, for the following reasons:

D2 discloses 5-HT-6 modulators, which are 4-sulfonamidoindole derivatives (see compounds of formula (le)(page 5 and Claim 6 of Do).

D4 disclose indoles which can be substituted with a sulfonamide moiety in any one of the positions 4-7 of the indole and which are useful to treat the same diseases as in the present case. Though D4 does not specifically disclose the role of 5HT-6 receptors, it clearly discloses the same application in terms of disorders to be treated. The possible discovery of a specific mechanism cannot be taken as an objective problem and eventually an inventive step as such.

The present general structures (Ia-b) differ from the compounds disclosed by D1 only in the positions of substitution with the ASO2N- and -(CH2)n-R1 moieties on the indole ring. D2 teaches that compounds substituted in position 4 with a sulfonamide function are useful as 5-HT-6 modulators. D3 discloses 5-HT-6 modulators, which can be substituted with a cycloalkyl moiety on the nitrogen of the indole ring (see e.g. examples 23, 28, 67 of D3) as for the present case. Having regard the minor modifications between the present compounds and the D1 compounds and in view of the D2-D4, the skilled person would have expected that the same qualitative effect would be maintain in such similar compounds. Moreover, the present R1 substituent can present very different chemical structures and therefore seems not to be relevant for retention of the claimed activity for the present compounds.

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The problem underlying the present application should thus be seen in providing of sulfonamide indole derivatives with unexpected or surprising effects compared to those of the closest prior art. An inventive step cannot be recognized as it is not yet shown by appropriate information, e.g. in form of experimental data, that substantially all the claimed compounds have un unexpected property or improved activity over the structurally closest prior art compounds (D1), which is attributable to the distinguishing feature of the invention.

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